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## Pregnancy outcome in South Australia

Verburg, Petra

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# CHAPTER 3

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## SEASONALITY OF GESTATIONAL DIABETES MELLITUS: A SOUTH AUSTRALIAN POPULATION STUDY

Petra E Verburg

Graeme Tucker

Wendy Scheil

Jan Jaap HM Erwich

Gus A Dekker

Claire T Roberts

# ABSTRACT

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## OBJECTIVE

To investigate whether there is a seasonal variation in the incidence of Gestational Diabetes Mellitus (GDM).

## RESEARCH DESIGN & METHODS

This retrospective cohort study of 60,306 eligible South Australian live born singletons during 2007-2011 recorded in the South Australian Perinatal Statistics Collection (SAPSC) examined the incidence of GDM in relation to estimated date of conception (eDoC). Fourier series analysis was used to model seasonal trends.

## RESULTS

During the study period, 3,632 (6.0%) women were diagnosed with GDM. Seasonal modeling showed a strong relation between GDM and eDoC ( $p < 0.001$ ). Unadjusted and adjusted models [adjusted for maternal age, body mass index (BMI), parity, ethnicity, socio-economic status and chronic hypertension] demonstrated the presence of a peak incidence occurring among pregnancies with eDoC in winter (June/July/August), with a trough for eDoC in summer (December/January/February). As this was a retrospective study, we could only use variables that had been collected as part of the routine registration system, the SAPSC.

## CONCLUSIONS

This study is the first population-based study to demonstrate a seasonal variation for GDM. Several maternal life style and psychosocial factors associated with seasonality and GDM may be influential in the pathophysiologic mechanisms of GDM. Ambient temperature, physical activity, nutrient intake and vitamin D levels may affect maternal physiology, fetal and placental development at the cellular level and contribute to the development of GDM. The mechanisms underlying these possible associations are not fully understood and warrant further investigation.

## Introduction

According to current guidelines, Gestational Diabetes Mellitus (GDM) is defined as carbohydrate intolerance, with onset or first recognition during pregnancy[1]. The incidence of GDM has increased several fold over the years from 1.9% in 1989-1990 to 4.2% in 2003-2004[2]. Currently, GDM is present in ~7% of all pregnancies, varying from 1% to 10% depending on the studied population[1]. The increase of GDM represents a significant economic burden on society, with direct medical costs of GDM in the United States in 2007 estimated at \$636 million[3].

GDM is an important contributor to maternal morbidity. Women with GDM have a 1.5-fold increased risk of hypertensive disorders of pregnancy[4], a 1.1-fold risk of delivery by elective caesarean section[5], a 7.4-fold risk of later Type 2 Diabetes (T2D)[6] and a 2.2-fold increased long-term risk of cardio-vascular disease (CVD)[7] compared to those women who had a normoglycaemic pregnancy.

GDM has also important offspring health effects, causing fetal morbidity and mortality. In GDM, the mother and the fetus are hyperglycaemic due to free transport of glucose through the placenta. The fetus is forced to increase its own insulin production[8]. Increased maternal glucose levels in pregnancy have been associated with increased frequencies of preterm birth, birth injury, a 5.0-fold increased risk of large for gestational age babies, a 2.2-fold increased risk of neonatal hypoglycaemia, requirement for intensive neonatal care and hyperbilirubinemia[5]. Pre-existing diabetes is has been associated with increased frequencies of congenital abnormalities[9]. In adulthood, offspring from mothers with GDM are at increased risk of developing obesity, impaired glucose tolerance, T2D and CVD[8].

The pathophysiology of GDM is not fully understood, but a number of risk factors have been implicated, such as maternal overweight and obesity, low maternal birth weight, race, increased maternal age, family history of T2D, history of previous fetal death and previous birth of a macrosomic infant[10,11]. Also, the recurrence rate of GDM in subsequent pregnancies has shown to be as high as 47.2%[12].

Other risk factors associated with GDM are nutrient intake, specifically high intake of saturated fat[10], high ambient temperature[13,14] and a low maternal vitamin D status[8,15–22]. Physical activity before and during pregnancy is associated with a significantly lower risk of developing GDM[23]. Since some of these risk factors have shown a seasonality[24–26], we hypothesized that the incidence of GDM may differ over the seasons. A seasonal trend of GDM could provide more insight on its pathophysiology. So far, only one study has reported seasonal changes in the prevalence of GDM[27].

In an effort to increase the knowledge of mechanisms regarding early pregnancy exposures that may influence the development of GDM, we aimed to assess the seasonal variation of GDM in a large population database. We focused on the estimated date of conception (eDoC) based on the hypothesis that the etiology of GDM lies early in pregnancy (and often pre-pregnancy), weeks to months before the clinical presentation of GDM.

## Research design and methods

### *Study design and setting*

This was a retrospective cohort study investigating the incidence of GDM in relation to eDoC in South Australia during 2007-2011. The majority of the state's population lives within the metropolitan area of the state capital, Adelaide. Adelaide (situated 34°S 138°E) is a coastal city with a Mediterranean climate, with long hot dry summers (December, January, February) and short cold rainy winters (June, July, August).

### *Database*

Data was sourced from the South Australian Perinatal Statistics Collection (SAPSC), maintained by the Pregnancy Outcome Unit (POU) of SA Health. The SAPSC collects information regarding the characteristics and outcome of all births, of any gestation, and stillbirths and terminations of pregnancy of at least 400 g birth weight or 20 weeks of gestation, in South Australia. The data were notified by hospital and homebirth midwives and neonatal nurses using a Supplementary Birth Record (SBR).

### *Data selection*

The study included live born singleton births in women with a known Body Mass Index (BMI) with a gestation of 20-42 weeks and a birth weight  $\geq 400$  grams. Terminations of pregnancy and women with pre-existing diabetes were excluded from the analyses.

When the clinician documented that the woman had GDM this was listed in the SAPSC. This was based on universal screening for GDM. All patients underwent a 50g glucose challenge test at 28 weeks. Patients with an abnormal glucose challenge test ( $\geq 7.8$  mmol/l) underwent a Glucose Tolerance Test (GTT); fasting  $\geq 5.5$  mmol/l and/or 2 hours value  $\geq 8$  mmol/l were the constant cut-off values to diagnose GDM in 2007-2011 in South Australia.

The eDoC was estimated based on the birth date and gestational age at birth. Gestational age was determined by best obstetric estimate and reported as completed weeks. This estimate was based on the dating ultrasound (performed at 8-13 weeks' gestation) supported by the known first day of the last menstrual period or by review of other ultrasonography. The database did not indicate how gestational age was specifically determined for each woman, but 99.1% of the women had an antenatal ultrasound.

### *Variables*

The variables included were maternal age, BMI, parity, ethnicity, socio-economic status (SES), civil status, smoking status at time of conception and in the second half of pregnancy, chronic hypertension, asthma and fetal sex. BMI was calculated before 20 weeks of gestation with the formula  $\text{BMI} = \text{weight in kg} / (\text{height in m})^2$ . Underweight was defined as  $< 18.5$  kg/m<sup>2</sup>, normal weight 18.5-24.99 kg/m<sup>2</sup>, overweight 25.00-29.99 kg/m<sup>2</sup>, obese 30.00-39.99 kg/m<sup>2</sup> and morbidly obese  $> 40.00$  kg/m<sup>2</sup>. The Australian Bureau of Statistics' Socioeconomic Index for Areas (SEIFA) population based quintile scores, derived from home address postcode, were used to indicate the SES[28]. For the civil status the participants were divided into those with a stable partner (married/de facto) and those without a stable partner.

### *Statistical analysis*

Data analysis was performed with SPSS version 21.0 (SPSS inc, 2013). Fisher's exact test and chi square test were used to compare categorical variables. Differences were considered significant when the  $p$ -value was less than 0.05.

In the primary analysis, eDoC was studied as a continuous variable using Fourier series methods (single cosinor analysis)[29]. Fourier series are considered to be the natural mathematic models for seasonality. To model the underlying seasonality of eDoC the first  $p$  pairs of term of the Fourier series was employed:

$$S(\theta_i, p) = \sum_{r=1}^p \{\beta_r \sin(r\theta_i) + \gamma_r \cos(r\theta_i)\}$$

In this series  $\theta_i$  is the point in the annual cycle that the  $i^{\text{th}}$  day on which eDoC occurred. Denoting the number of days between 1 January 1950 and the  $i^{\text{th}}$  day of woman's eDoC as  $D_i$ , we calculated this angle in radians thus:  $\theta_i = 2\pi (D_i \bmod 365.25) / 365.25$ . Thus, seasonal effect of the eDoC on the binary pregnancy outcome is modelled by adding  $S(\theta_i, p)$  to the linear predictor of a logistic regression model so that  $\beta_i$  and  $\gamma_i$  become parameters in a simple linear model. In these data, the first pair of Fourier terms (F1 model: sine and cosine) was significant based on a likelihood ratio test ( $\alpha = 0.10$ ), allowing us to use these in this study.

Seasonality of GDM was initially investigated using a univariate logistic regression analysis using eDoC divided in months as predictor. Subsequently, the model was run with and without adjustment for potential confounders defined *a priori* (maternal age, BMI, parity, ethnicity, SES, civil status, smoking status, chronic hypertension, asthma and fetal sex) using effects coding. The average of all months together was the reference in the model when using effects coding.

Subsequently, multivariate logistic regression models (Fourier series analysis) were fit by specifying a full model with all available data on potential modifiers and confounding variables defined *a priori* (see above). Covariates were considered not to be influential and were removed from the model via backward elimination if their contribution to the model was not significant ( $p > 0.05$ ). Coefficient correlation tests were performed after every step of the model fitting process to test the goodness of fit.

### Ethics

The existence of personal identifying information in the SAPSC was eliminated from the data file for analysis to ensure that confidentiality of all patient records was maintained. The study protocol was approved by the Human Research Ethics Committee of the South Australian Department of Health [HREC/13/SAH/97].

## Results

### Study population

Of a total of 99 973 births registered between January 2007 and December 2011, 60 306 (60.3%) births were eligible for the study. We excluded 25 (0.03%) births with a gestational age of <20 weeks or >42 weeks, 269 (0.27%) births in which the neonate had a birth weight <400 grams, 121 (0.12%) terminations of pregnancy because of chromosomal and/or congenital anomalies, 357 (0.36%) stillbirths, 2 961 (2.96%) non-singleton births, 35 542 (35.55%) births with inadequately recorded maternal BMI and 392 (0.39%) births in which pregnancy was complicated by pre-existing diabetes.

### Incidence of GDM

During the study period, a total of 3 632 (6.0%) women were diagnosed with GDM. The incidence of GDM increased significantly over the years from 4.9% in 2007 to 7.2% in 2011 ( $p < 0.001$  (Table 1)). The characteristics of the study population are presented in Table 2. Compared to non-GDM pregnancies, risk factors for GDM included age >30 years, multiparity, increased BMI, Asian ethnicity, low and very low SES and a history of chronic hypertension.

Table 1. Incidence of GDM 2007-2011.

	Incidence GDM complicated births		All births
Year of birth	N	%	N
2007	396	4.9	8,001
2008	600	5.4	11,062
2009	780	6.0	12,947
2010	795	5.8	13,633
2011	1,061	7.2	14,663

GDM, gestational diabetes mellitus

Table 2. Characteristics of the study population.

Characteristic	GDM complicated births		Non-GDM births		All births	
	n	%	n	%	n	%
<b>Age (in years)</b>						
< 20	45	1.2	2,653	4.7	2,698	4.5
20-25	288	7.9	9,941	17.5	10,229	17.0
25-30	876	24.1	17,593	31.0	18,469	30.6
30-35	1,265	34.8	16,480	29.1	17,745	29.4
35-40	876	24.1	8,298	14.6	9,174	15.2
> 40	281	7.7	1,710	3.0	1,991	3.3
<b>BMI</b>						
< 18.5 Underweight	61	1.7	1,837	3.2	1,898	3.1
18.5-25 Normal weight	1,091	30.0	27,022	47.7	28,113	46.6
25-30 Overweight	923	25.4	15,217	26.8	16,140	26.8
30-40 Obese	1,155	31.8	10,675	18.8	11,830	19.6
> 40 Morbid obese	401	11.0	1,924	3.4	2,325	3.9
<b>Parity</b>						
Nulliparous	1,452	40.0	24,775	43.7	26,227	43.5
Multiparous	2,179	60.0	31,900	56.3	34,079	56.5
<b>Ethnicity</b>						
Caucasian	986	27.2	9,069	16.0	10,055	16.7
Aboriginal or TSI*	94	2.6	1,426	2.5	1,520	2.5
Asian	695	19.1	5,186	9.2	5,881	9.8
Other	197	5.4	2,457	4.3	2,654	4.4
<b>Socio-Economic Status</b>						
Very low (0-20%)	1,074	29.6	14,660	25.9	15,734	26.1
Low (20-40%)	786	21.6	12,020	21.2	12,806	21.2
Medium (40-60%)	695	19.1	11,811	20.8	12,506	20.7
High (60-80%)	549	15.1	9,352	16.5	9,901	16.4
Very high (<80-100%)	505	13.9	8,621	15.2	9,126	15.1
<b>Civil status</b>						
With stable partner	3,296	90.8	49,905	88.1	53,201	88.2
Without stable partner	335	9.2	6,770	11.9	7,105	11.8
<b>Smoking</b>						
at time of conception†	633	17.4	11,314	20.0	11,947	19.8
2nd half of pregnancy‡	403	11.1	7,780	13.7	8,183	13.6
<b>Medical complications</b>						
Chronic hypertension	121	3.33	546	0.96	667	1.11
Asthma	270	7.44	3,643	6.43	3,913	6.49
<b>Fetal sex</b>						
Male	1,737	47.84	27,631	48.75	29,368	48.70
Female	1,895	52.19	29,044	51.25	30,939	51.30

\* Torres Strait Islander

† Analysis of 59,881 subjects (4 243 smoking vs. 55 638 non-smoking)

‡ Analysis of 59,411 subjects (4 213 smokers vs. 55 198 non-smokers)

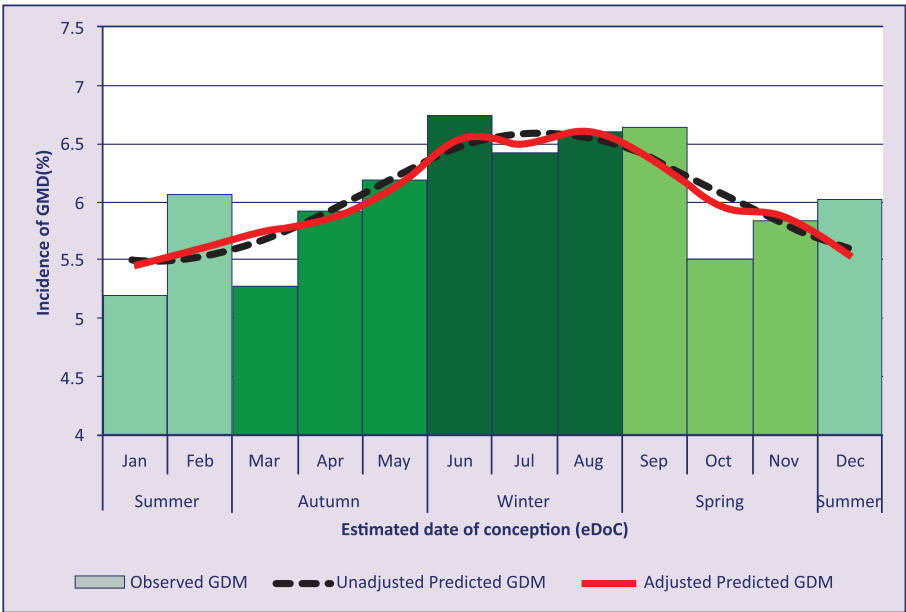


*Seasonal models of GDM*

The observed, unadjusted and adjusted probabilities for GDM by eDoC are shown in Figure 1. Both unadjusted and adjusted predicted probabilities were based on the logistic regression model fitting the first pair of Fourier series terms and showed that GDM was significantly related to season of eDoC ( $p < 0.001$ ).

The highest incidence of GDM was observed among pregnancies with eDoC in June (6.74%). Unadjusted seasonal modelling showed peak incidence of GDM for pregnancies with eDoC in July (6.55%), while adjusted seasonal modelling showed a peak incidence of GDM for pregnancies with eDoC in August (6.60%).

The lowest incidence of GDM was observed among pregnancies with eDoC in January (5.17%). Seasonal modelling showed a corresponding pattern. Both unadjusted and adjusted predicted incidence showed a nadir for pregnancies with eDoC in January (5.46% and 5.41%, respectively).



**Figure 1. Fitted seasonality for GDM by estimated date of conception (eDoC).** The histogram represents the observed probability of GDM by eDoC (divided in calendar months). The dashed line represents the unadjusted Fourier fit for the incidence of GDM by eDoC. The bold line represents the adjusted Fourier fit for the incidence of GDM by eDoC. The adjusted Fourier fit was adjusted for: maternal age, BMI, parity, ethnicity, socio-economic status and chronic hypertension.

## Conclusions

### *Increasing incidence of GDM*

The incidence of GDM in South Australia increased significantly from 4.9% in 2007 to 7.2% in 2011. This is consistent with the trend in the United States, where incidence increased from 1.9% in 1989-1990 to 4.2% in 2003-2004[2]. This increase in incidence is not surprising giving the parallel rise in prevalence of T2D and obesity[8] and also increased focus on detection of GDM in pregnancy in recent decades.

### *Seasonality of GDM*

In this large South Australian birth cohort, we observed a clear seasonality in the incidence of GDM based on eDoC. The peak incidence was observed for pregnancies with eDoC during winter and the lowest incidence for pregnancies with eDoC during summer.

So far, only one study reported a seasonal pattern in the prevalence of GDM[27]. The study, carried out in Wollongong (coastal city, 34 °S), Australia, in 2010-2014, studied the results of 7,343 oral glucose tolerance tests performed between 24 and 28 weeks' gestation and a total of 516 (7.0%) women were diagnosed with GDM. The study found that in winter the median 1-hour- and 2-hour glucose results were significantly lower compared to the overall 1-hour- and 2-hour results. A post-hoc analysis indicated that the prevalence of GDM based on the 1-hour- and 2-hour results was higher in summer (respectively 4.6% and 4.4%) and lower in winter (respectively 2.6% and 2.4%) compared to the overall prevalence of GDM. Interestingly, they did not find a seasonal variation in fasting glucose levels. Diagnosis in summer corresponds with eDoC in winter, which concurs with our data.

Two previous studies reported that there was no seasonal pattern in the incidence of GDM[30,31]. An older and small study, carried out by the same group in Wollongong, Australia, in 1993-1994, studied a total of 2,749 subjects with an overall incidence of GDM of 6.7% (183 subjects). The seasonal fluctuation of GDM showed an incidence of 7.4% (5.6-9.2) in autumn (March to May), corresponding with a conception in late winter/early spring (August to October). An incidence of 5.5% (3.5-7.5) was observed in winter, corresponding with a conception in late spring/early summer (November to January). However, this seasonal fluctuation was not statistically significant[30].

Another small study was conducted in Plymouth (coastal city, 50 °N), United Kingdom, in 1996-1997. A total of 4,942 pregnant women were studied, with an overall prevalence of GDM of 1.8% (90 women). The incidence of GDM ranged from 2.3% (1.5-3.2) in spring, corresponding with conception in autumn and the lowest incidence of GDM of 1.4% (0.8-2.3) in winter, corresponding with conception in summer. However, this seasonal fluctuation was also not statistically significant[31].

Small sample size (183 and 90 women with GDM, respectively) and therefore lack of statistical power is the most likely reason these two prior small studies could not detect significant seasonality. Also, Plymouth has a very mild climate, with little fluctuation in temperature and sun exposure and therefore seasonality is expected to be less distinct than in more extreme climates. The actual trends were similar to the findings in this large South Australian cohort. Inconsistencies in the magnitude of GDM incidence is not surprising in this type of study, since population composition, environmental exposures and climates differ in different regions of the world.

### *Seasonality of Type 2 Diabetes*

Previous studies have analysed the seasonal variation in T2D. A Hungarian study, investigating 26,695 cases of T2D from 1999-2004 in Csongrad County, Hungary (46 °N), demonstrated a sinusoidal pattern for the incidence

of T2D in both males and females, with a peak incidence in Spring and a trough in Summer[32]. This finding in T2D diagnosis is not in line with what we found in our GDM data. However, there are multiple potential reasons for this apparent discrepancy. Standardized testing in pregnancy between 24-28 weeks' gestation eliminates patient's and doctor's delay in diagnosing GDM, versus T2D. Also, even though T2D and GDM are associated, the timelines to diagnosis are quite different.

Considering the Developmental Origins of Health and Disease concept[33,34], suggesting prenatal programming of physiology, it can be hypothesized that T2D might show a seasonal variation depending on month of birth of the individual. Several studies have investigated this relationship with conflicting results[35,36]. An Ukrainian study in three regions: Charnigov (51 °N), Kherson (46 °N) and Rivne (50 °N) of 17,510 male and 34,704 female type 2 diabetic patients showed a seasonal pattern with a peak in births in April for both males and females, and a nadir of births for males in November and for females in December[35]. However, a large Danish cohort study, investigating 223,099 people, of whom 12,468 developed T2D, did not find a relationship between season of birth and the incidence of T2D[36]. However, as this study used time of birth to indicate season it did not account for the fact that there was a range of gestational ages from preterm to term, possibly leading to confounding.

#### *Possible mechanisms underlying the seasonality of GDM*

In our analyses we observed a seasonal variation of GDM after adjusting for maternal age, BMI, parity, ethnicity, SES and chronic hypertension. These characteristics change relatively little from month to month and are not or barely modifiable. Owing to the retrospective character of the present study we could not adjust for several other meteorological, maternal life style and psychosocial factors associated with seasonality, such as ambient temperature, physical activity[24], nutrient intake[25] and vitamin D status[26]. Several studies have shown that these factors have been associated with the risk of GDM[13,21,23,37,38].

#### *Meteorological factors*

Studies suggest that the incidence of GDM can be associated with variable seasonal meteorological factors, specifically ambient temperature. A study in Porto Alegre (temperate climate 30°S), Brazil, of 1030 pregnant individuals compared serum glucose in glucose tolerance tests performed between 24 and 28 weeks' gestation with ambient temperature. For every degree increase in ambient temperature, serum glucose increased by 0.07 mmol/L[13]. The Brazilian study suggests that the incidence of GDM may be highest in summer. GDM is generally diagnosed between 24 and 28 weeks of gestation[39], which corresponds with conception in winter. This is consistent with our findings that the highest incidence was seen when eDoC occurred in winter.

#### *Physical activity*

A systematic review and meta-analysis of 34,929 participants and 2,813 cases of GDM showed that pre-pregnancy and early pregnancy physical activity were significantly protective against GDM (55% and 24%, respectively, lower risk of GDM)[23]. During pregnancy, the body is subject to metabolic changes, especially during the third trimester, when GDM is typically diagnosed. GDM is thought to reflect an impaired capacity of the woman to adjust to these metabolic changes, due to relative  $\beta$ -cell dysfunction, i.e. insulin secretion not keeping up with the exaggerated state of insulin resistance, resulting in abnormal glucose tolerance. An inverse association is plausible and it can be suggested that pre-pregnancy and early pregnancy exercise may improve overall fitness and glucose tolerance[23].

A systematic review has shown seasonal variation in the levels of physical activity in non-pregnant study populations. In general, the highest levels of physical activity were seen in spring and summer, while the lowest levels were seen in winter[24].

These results are also consistent with our findings that the lowest incidence of GDM occurred with summer conceptions when women were more likely to be physically active. However, we know of no studies reporting seasonal variation in levels of activity in pregnancy.

#### *Nutrient intake*

Seasonal variation of nutrient intake may affect metabolism and this may also provide clues to understand the seasonal variation of GDM. Prospective[25] and retrospective[40] studies have demonstrated a seasonal trend in nutrient intake in pregnant women. Unfortunately, data are not completely consistent. Previous studies have shown that a high total fat intake at time of diagnosis, in most cases at 24-28 weeks' gestation, is associated with an increased risk for GDM[10,37]. Interestingly, the total fat intake in one study was shown to be highest in Spring[25], whereas the other showed the highest fat intake in Summer[40]. The latter corresponds to conception in autumn and winter, which is consistent with our findings.

#### *Vitamin D*

Serum vitamin 25(OH)D<sub>3</sub> levels have a seasonal variation, with highest concentrations in summer and autumn and lowest concentrations in winter and spring[26]. This is not surprising, since vitamin D is mainly derived from skin exposure to sunlight and dietary intake[41]. The lowest dietary intake of vitamin D was seen in autumn and the highest in spring[25,40].

A recent meta-analysis of studies published between 2008 and 2015 with over 9200 women showed that low vitamin D status associates with GDM[42]. A vitamin D response element in the human insulin receptor gene promoter has been identified[43], suggesting a potential role for vitamin D in enhancing insulin secretion. Wang et al. demonstrated that higher vitamin D levels stimulate increased  $\beta$ -cell mass and as such potentially also  $\beta$ -cell insulin secretion capacity[44]. However, other studies have reported conflicting results. The association between vitamin D and its specific role in the development of GDM is contentious and merits further investigation.

#### *Strengths and weaknesses of the study*

A major strength of this population-based study lies in the large number of analysed births. The SAPSC collects information regarding the characteristics and outcome of all births in SA, notified by hospital and homebirth midwives and neonatal nurses.

As this was a retrospective study, we could only use variables that were already listed on the dataset. GDM was listed in the SAPSC when the clinician documented that the women had GDM. In the study period there were stable criteria for diagnosis and a consistent approach used by all care providers. This study included only the live born babies during the study period. However, third trimester stillbirth is not very common, so it is unlikely that this has influenced the seasonal variation of GDM observed in this study. Since 2007 SAPSC began collecting data on BMI of the pregnant women. We used BMI as one of our inclusion criteria, which meant we had to exclude 35.55% of the total births between January 2007 and December 2011. SAPSC validation studies have shown that notifications of births in SA made by hospital and homebirth midwives and hospital neonatal nurses on the SBR were robust for the parameters studied[45]. The pregnancies analysed in this study could therefore be considered as a proper representation of the South Australian and Australian population.

### Conclusion and further research

This representative study has identified a statistically significant seasonal variation in the incidence of GDM. The etiology of GDM involves exposures that occur before it is clinically recognised. This suggests that the pathophysiologic mechanisms explaining (the seasonality of) GDM may have their origin early in pregnancy. These epidemiological data also support exploring the contributions of exposures with seasonal periodicity to GDM, such as environmental and life style differences.

The effect size of seasonality of GDM presented in this paper is statistically significant, but unlikely to have direct clinical implications. However, our data point to specific biological mechanism(s) that warrant further investigation that may provide potential targets for the prevention of GDM. In particular, exposure(s) early in pregnancy deserve attention. These may include season of conception and vitamin D.

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